Barotropic tidal and wind-driven larval transport in the vicinity of a barrier island inlet

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ABSTRACT

Fish species such as Atlantic menhaden (Brevoortia tyrannus) must often transit relatively small barrier island inlets to move from their continental shelf spawning grounds to estuarine nurseries, where they spend their juvenile phase. Physical transport through these inlets is strongly influenced by tides, winds, local geometry and bathymetry, and can be very different from that occurring on the shelf. In March 1996, an extensive multidisciplinary field experiment was conducted to identify larval transport pathways in the vicinity of Beaufort Inlet, NC (Blanton et al., 1999; Churchill et al., 1999; Forward et al., 1999). One of the most interesting results from the field study was significant spatial inhomogeneity in the larval catches (Forward et al., 1999).

This paper describes a computer modelling study of tidal and wind-driven circulation and accompanying larval transport characteristics in the vicinity of Beaufort Inlet. Primary conclusions are as follows: (i) tidal currents are quite effective at creating net larval transport into the inlet; (ii) in the absence of wind forcing, to explain the spatial distribution of larval catches observed in the field experiment, the primary source of larvae must have been to the east of the inlet; (iii) the spring/neap cycle accounts for a variation of approximately 40% in larvae ingress; (iv) only a few wind directions enhance larvae ingress over the case of purely tidal forcing—wind blowing toward the north enhances ingress at the surface, and winds blowing toward the south and the east enhance ingress at the bottom; (v) if larvae are evenly distributed in space outside the inlet, then strong east wind (such as 10 m s\(^{-1}\)) can cause larvae to move through the inlet in a way that is consistent with the larval catches in the field experiment. However, local wind records revealed that no significant eastward-blowing winds occurred during or immediately preceding the field experiment.

Key words: larval transport, inlet studies, modelling, circulation

INTRODUCTION

The life cycle of fish species such as Atlantic menhaden (Brevoortia tyrannus) that spawn in continental shelf waters and spend their juvenile phase in estuarine nursery grounds depends critically on physical processes that transport larval fish from spawning areas to juvenile nurseries (Miller et al., 1984). In the South Atlantic Bight, transport on the mid-to inner continental shelf is primarily wind driven (Werner et al., 1999), with additional influences from Gulf Stream intrusions (Stegmann et al., 1999). However, in most cases, to reach estuarine nursery areas, larvae must find and transit relatively small barrier island inlets. Physical transport through these inlets is strongly influenced by tides, local geometry and bathymetry, so is very different from that occurring on the shelf. Thus, a dedicated effort to understand this phase of larval transport is critical. In addition, since many inlets are modified by dredging and jetty structures to maintain navigability, it is important to understand inlet transport as a basis for assessing the implications of these actions.

Barotropic tidal circulation patterns on the seaward side of an idealized, symmetric inlet are reasonably well established (see, for example, Zimmerman, 1981; Mehta, 1988; Oertel, 1988). On ebb tide, a boundary-separated jet forms that preferentially transports water out of the estuary in line with the inlet centre. In contrast, on flood tide, the inflow is uniformly distributed in a convergent pattern toward the inlet.

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The difference between the ebb and flood patterns sets up a steady tidal residual circulation that is offshore in the inlet centre, towards the inlet near shore, and contains counterclockwise eddies on either side of the offshore flow. (Looking in the offshore direction from the inlet mouth, these eddies rotate clockwise to the left of the inlet and clockwise to the right of the inlet.) In the Northern Hemisphere, the clockwise eddy is generally stronger than the counterclockwise eddy, owing to the influence of the Earth’s rotation. The strength of both eddies is enhanced by offshore sloping topography (Chao, 1990). On the landward side of an inlet, marsh systems, shoals and/or the shoreline often prevent a similar pattern from forming and largely dominate the circulation patterns that occur there.

Baroclinic effects associated with fresh-water discharge have also been studied in idealized inlets. Fresh-water discharge superimposes estuarine-like circulation (seaward flowing buoyant water overriding shoreward flowing dense water) over the barotropic tidal circulation. During ebb tide, buoyant water flows over denser shelf water, creating a stably stratified water column and a plume outside of the inlet. Owing to the Earth’s rotation, in the Northern Hemisphere, plumes are characterized by clockwise circulation that enhances the strength of the clockwise residual eddy and diminishes the strength of the counterclockwise residual eddy near the surface (Chao, 1990; Kapolnai et al., 1996; Wheless and Valle-Levinson, 1996). A weaker counterclockwise circulation develops under the plume and a buoyant surface coastal current that moves away from the inlet to the right (in the Northern Hemisphere) may also form (Chao, 1990). During flood tide, stratification is largely destroyed as denser shelf water is transported through the inlet (Wheless and Valle-Levinson, 1996).

Past inferences about larval transport through tidal inlets and adjacent estuaries have been drawn from the relationship between the vertical position of larvae in the water column and the phase of the tidal cycle (Hill, 1991; Smith and Stoner, 1993; DeVries et al., 1994; Chen et al., 1997; Power, 1997), considerations of subtidal physical processes, such as buoyancy-induced circulation or gradients in wind-driven water level set up (Pietrafesa and Janowitz, 1988), and the potential for larvae to remain entrapped in gyres or other areas of slow flow within an estuary (Føre and Baxter, 1972; Trinast, 1975). However, there has been little work on net larval transport associated with tidal dispersion, as described by Zimmerman (1976), which results from the interaction between a spatially varying oscillatory current (such as a tide near irregular bathymetry) and a spatially varying residual current (such as a tidal or wind-driven residual). This mechanism can account for significant net transport when geometric and topographic variability is large over spatial scales less than a tidal excursion length (Geyer and Signell, 1992), as can be expected near tidal inlets. Perhaps the main reason why tidal dispersion has not been widely considered in larval studies is because it requires detailed information about the horizontal structure of the flow. This information is most readily obtained from numerical circulation models (see, for example, Seaberg, 1988; Geyer and Signell, 1992), the development of which can be quite challenging in the highly complex geometry and topography that typically surround a tidal inlet.

In the present study, we consider larval transport in the vicinity of Beaufort Inlet, North Carolina (Fig. 1). The inlet is approximately 1 km wide and connects a complicated estuarine system to Onslow Bay and the South Atlantic Bight. Immediately inside the inlet,
flow is directed through the Morehead Channel, the Radio Island Channel or the Shackleford Channel. Habitat quality is dramatically different in the waters upstream of these channels, as the Morehead Channel connects to the port of Morehead City, the highly developed Bogue Sound and the relatively undeveloped Newport River Estuary; the Radio Island Channel connects to the fully developed waterfront of Beaufort, NC, and the Newport River Estuary; the Shackleford Channel connects to relatively pristine, high-quality nursery areas.

Flow near the inlet is dominated by the semidiurnal, $M_2$ tide with additional episodic contributions from wind and fresh-water forcing (Klavans, 1983; Logan, 1995). Beaufort Inlet was the site of a multidisciplinary, larval transport field experiment conducted during two consecutive neap tides in March 1996. Physical data from this experiment (Blanton et al., 1999) indicate that fresh-water discharge was significant during the first period (salinity ranged from 20 to 34 psu over the tidal cycle) but much less during the second period (salinity ranged from 32.5 to 34.5 psu over the tidal cycle). During both periods, the water column was vertically well-mixed both in the inlet mouth and in the channels shoreward of the inlet, through virtually the entire tidal cycle. Despite the difference in fresh-water discharge, patterns of larval catches inside the inlet remained remarkably consistent between the two periods (Forward et al., 1999). The spatial scales of the inlet, the presence of a well-mixed water column and the apparent insensitivity of larval catches to buoyant discharge encouraged us to focus initial modelling efforts on the barotropic component of the tidal and wind-driven circulation. Specifically, we have sought to (i) understand basic tidal circulation and tidal dispersion in the vicinity of Beaufort Inlet; (ii) to provide insight into the field observation that an order of magnitude more larvae enter the inlet and move eastward through the Shackleford Channel than move westward through the dredged Morehead Channel (Forward et al., 1999); and (iii) to assess the role of wind forcing vs. tidal forcing in determining larvae ingress through Beaufort Inlet.

**BEAUFORT INLET CIRCULATION**

Barotropic circulation in Beaufort Inlet was computed using two complementary numerical hydrodynamic models. Tidal circulation was computed using the depth-integrated version of the ADCIRC model (Luebth et al., 1992), while wind-driven circulation was computed in three dimensions using the FUNDY5 model (Lynch et al., 1992).

Both ADCIRC and FUNDY5 utilize the finite element method on linear triangles for their spatial discretization and have been run on a grid (Figs 2 and 3) that stretches from approximately the North Carolina–Virginia border to the South Carolina–Georgia border. The grid extends from the deep ocean (more than 5000 m deep) to estuarine tidal flats that wetted and dried (in the tidal simulations only), and has element sizes ranging from approximately 25 km in deep water to 25 m near shore.

ADCIRC is a fully nonlinear, time-stepping model that accurately generates overides and compound tides in strongly nonlinear tidal flows (Grenier et al., 1995). The two-dimensional assumption is appropriate for energetic tidal flows where the water column remains well mixed and the water column depth is small compared with the Ekman depth. However, a two-dimensional model is not appropriate for wind-driven flows in shallow enclosed or semi-enclosed areas, where

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**Figure 2.** Finite element grid used for the Beaufort Inlet modelling study.
the presence of land can force an upwind circulation at depth that opposes the generally down-wind circulation at the water surface. Therefore, the FUNDY5 model, which provides a harmonic in time solution to the linearized, three-dimensional equations of motion, was used to describe the wind-driven current fields.

Tidal model results
The ADCIRC model was forced using the five primary astronomical constituents: $M_2$, $S_2$, $N_2$, $K_1$ and $O_1$. Forcing was applied throughout the domain using the direct tide producing tidal potential terms, and along open boundaries using elevations extracted from the FES95.2 global tidal database (Le Provost et al., 1998). A hybrid bottom friction formulation was used that approached a quadratic relationship with a friction coefficient of $0.0025$ in water deeper than 1 m, and a Manning relationship (Manning’s $n = 0.016$) in shallower water. Based on preliminary model runs, a spatially constant lateral viscosity of $2 \text{ m}^2 \text{s}^{-1}$ was used to maintain numerical stability. While some noise appears in the residual velocity field at a few locations, this was considered to be preferable to the reduction in lateral shear that resulted from using a significantly larger value. The model time step (Courant number limited) was 5 s. The model was spun-up from rest for 10 days, and the subsequent 30 days were harmonically analysed at all grid points within the domain for the five astronomical constituents as well as for the steady residual component and 24 nonlinearly generated compound tides and overides. As described in the following, the model results were in good agreement with available observational data, so no calibration of the model was attempted (such as by adjusting the open boundary conditions, water depths or bottom friction coefficients).

The amplitude and Greenwich phase of the dominant semidiurnal ($M_2$) and diurnal ($K_1$) tidal constituents are shown in Fig. 4. The semidiurnal tide on the shelf is a co-oscillation between the shelf waters and the Atlantic Ocean, and arrives nearly in phase along much of the US east coast. On entering the inlet, it is strongly influenced by bottom friction, owing to the shallow water depths and high current velocities, and is quickly attenuated (Fig. 4(a)) and slowed (Fig. 4(b)) as it propagates into the sounds. The diurnal tide on the shelf is a southward propagating wave that attenuates more slowly inside the estuary (Figs 4(c) and 4(d)), owing to its lower velocity and the resulting reduced frictional influence (Grenier et al., 1995).

Modelled vertically integrated tidal velocities (the full model response to the five astronomical constituent forcing) for the area around Beaufort Inlet are shown at four phases of the tide in Fig. 5. Velocities reach approximately 1 m s$^{-1}$ at maximum flood and ebb in the inlet throat (Figs 5(b) and 5(d)). The ebb flow exits the inlet with a jet-like structure, while the flood flow is fairly uniformly distributed in space outside the inlet. It is clear that any similar pattern inside the inlet is disrupted by the islands and shoreline. Near slack before ebb (Fig. 5(a)) and slack before flood (Fig. 5(c)), the tide can be seen to turn in the Morehead Channel (west of the inlet) earlier than it turns in the Shackleford Channel (east of the inlet). The model indicates a difference of approximately 45 min between the time of slack tide in these two channels, which corresponds closely to differences in the time of slack tide measured during the larval transport field experiments conducted during March 1996 (Blanton et al., 1999; Churchill et al., 1999). The velocity field at slack before flood (Fig. 5(c)) also suggests that initial flooding of the inlet consists of ‘new’ ocean water travelling west along Shackleford Banks entering the eastern side of
Figure 4. Tidal elevation amplitude (metres) and Greenwich phase (degrees) of the largest semidiurnal ($M_2$) and diurnal ($K_1$) tidal constituents for the area around Beaufort Inlet.

the inlet, and ‘old’ estuary water from the previous ebb plume that is re-entrained into the inlet along the western side. This corresponds closely to observations that, at the onset of flood tide, ocean water enters the estuary in roughly the eastern third of the inlet and is visually very distinct from water in the remainder of the inlet (R. Lueitich, personal observation).

The vertically integrated Eulerian tidal residual velocity field generated by the model is shown in Fig. 6. This steady circulation, with maximum velocities of 15 cm s$^{-1}$, indicates the existence of offshore flow in the inlet channel and near-shore flow toward the inlet, as suggested by idealized barotropic inlet studies. However, the absence of counterrotating eddies on either side of the offshore flow represents a significant departure from these earlier studies. Instead, one large recirculation zone, approximately 4 km in diameter, appears to the west of this flow. At this point, we are pursuing additional studies to verify the existence of this feature and to identify factors (such as the angle of the inlet channel with respect to the coastline, the difference in the ebb tidal deltas on either side of the inlet) that may control its size and strength. The structure of the residual field suggests that a plume exiting the inlet would tend to move offshore, to the west, and might be re-entrained along the western side of the inlet, while offshore ocean water would be entrained along the eastern side. In the inlet throat, the residual flow is offshore on the western side of the inlet and onshore on the eastern side of the inlet. This is in good agreement with acoustic Doppler current profiler measurements made by Churchill et al. (1999) and current meter mooring data collected by Logan (1995). Inside the inlet, there appears to be a residual velocity directed from Shackleford Channel to Morehead Channel.

Model results were verified against water surface elevation and velocity data collected in the 1970s by the National Ocean Survey (Klavans, 1983). Observed tidal elevations at 16 stations in the vicinity of Beaufort Inlet (Fig. 7(a)) are compared with model results in Table 1. Both amplitude and phase are well simulated (typically within 10% amplitude and 10° phase) for the
Figure 5. Vertically integrated tidal velocities for the area around Beaufort Inlet: (a) slack before ebb; (b) maximum ebb; (c) slack before flood; (d) maximum flood.

primary astronomical constituents. The $M_2$ phase is typically 10–20 min (5–10°) early in the model and this probably results from slight errors in the offshore boundary forcing. Other consistent differences between the model and the observational data occur at locations in small channels (such as stations 4 and 18) or furthest from the inlet (such as stations 5, 10 and 15). This is probably because of coarseness of the model grid (relative to some of the smaller bathymetric and geometric scales) and inaccurate bathymetry (which can shift significantly yet is rarely surveyed outside of the shipping channel). The $M_4$ overtide is not as well represented as the astronomical constituents; rather, this nonlinearly generated constituent is typically too large in the model. Generation of $M_4$ is strongly dependent on water depth (Grenier et al., 1995) and geometric irregularities, suggesting that inaccurate bathymetry and shoreline representations account for a significant part of this problem. There is also some question about the fidelity of the observational data, since some closely spaced stations have very large differences in the observed $M_4$ component (such as stations 11 and 12).

Tidal velocity data are available for 10 positions near the inlet (Fig. 7(b)). Figure 8 presents comparisons between the modelled, depth-averaged tidal ellipses and observed tidal ellipses for the $M_2$ and $K_1$ tides. Flow is highly rectilinear and directed by the channel geometry. The model tends to underpredict the observed velocity magnitude in the inlet mouth (stations 6 and 7) by about 15% but reasonably reproduces the remaining velocity observations. Because of rapid bathymetric and geometric changes, flow near the inlet mouth is subject to strong horizontal shear. Therefore, these discrepancies may be the result of small spatial errors in current meter positions relative to the model.
Figure 6. Vertically integrated, Eulerian tidal residual velocities near Beaufort Inlet.

Figure 7. Locations of tidal elevation and velocity stations where observational data are available from a study by the National Ocean Survey (Klavans, 1983). Station numbering is from Klavans (1983).

domain. The model has a sufficiently accurate representation of the channels in the vicinity of the inlet that the directions are well reproduced except at stations 9 and 11. These two stations are far up into the estuary, where model bathymetry or geometry presumably does not reflect local conditions.

Additional model verification was performed by comparing model fluxes through the three primary channels.
Table 1. Comparison between modelled tidal elevation amplitudes and phases and NOS observational data (Klavans, 1983). Station numbering is from Klavans.

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NOS, National Ocean Survey.

inside the inlet with fluxes measured using a shipboard acoustic Doppler current profiler. These data were obtained from repeated passes across the channels during the second larval transport field experiment conducted in March 1996 (Churchill et al., 1999). Model fluxes were computed by integrating the model results along lines matching the field tracks. The model underpredicts maximum flux through the small Radio Island Channel by as much as 50%, but is within about 15% of the observed flux through the main Morehead and Shackleford Channels (Fig. 9). In each channel, the model captures both the phasing and the diurnal inequality quite well.
Wind model results
While tidal currents dominate the instantaneous flow near Beaufort Inlet and tidal dispersion can account for considerable net larval transport (see below), significant energy also exists at subtidal frequencies (Blanton et al., 1999). In the presence of dominant tidal circulation, meteorologically forced subtidal circulation should be compared with the tidal residual circulation and with the tidal current shear, to determine the potential of the former to modify net transport as a result of tidal dispersion. We considered this effect by generating steady-state, three-dimensional circulation fields using the FUNDY5 model, driven by steady and spatially uniform wind fields that are climatologically representative of spring weather near Beaufort Inlet. Although wind events are clearly transient in nature, the shallow depths near the inlet suggest that the system will spin-up rapidly as it approaches steady state. Therefore, this limit is useful for an initial understanding of the role of wind in near-inlet circulation.

Since FUNDY5 is a linear model, it was necessary to develop a linear bottom friction coefficient and a constant-in-time vertical eddy viscosity that reflected turbulence generated via bottom mixing by the strong, tidally dominated currents. We computed a linear bottom friction coefficient $k_i$ as

$$k_i = \frac{c_u |U|}{h}$$  \hspace{1cm} (1)

where $c_u$ is the quadratic drag coefficient ($= 0.0025$), $h$
Figure 9. Comparisons between tidal fluxes computed by the model with those measured using a shipboard acoustic Doppler current profiler during the March 1996 field experiments (Churchill et al., 1999).

\[
E_z = 0.1hU^*
\]  \hspace{1cm} (2)

where \(U^*\) is the friction velocity. Both \(1U1\) and \(U^*\) were taken as spatially varying root mean square values determined from the ADCIRC solution. Minimum values were set, requiring \(k_1 > 0.00025\) s\(^{-1}\) if \(h > 50\) m and \(E_z > 0.0005\) m\(^2\) s\(^{-1}\).

During the winter–spring period, the climatological mean wind speed as measured by an NOAA buoy near Cape Lookout, NC, is approximately 6 m s\(^{-1}\) with a standard deviation of 2.5 m s\(^{-1}\). The direction toward which the wind blows ranges from north-east to south-east, with an average direction being toward the east during March (Werner et al., 1999). We illustrate basic characteristics of the wind-driven circulation near Beaufort Inlet by presenting the modelled current field at the surface and at the lowest grid point (essentially 1 m above the bottom) for a wind of 7.5 m s\(^{-1}\) directed toward the east (Figs 10(a) and 10(b)) and directed toward the north (Figs 10(c) and 10(d)).

Because of the shallow water depths, the surface Ekman layer is depth-limited and eastward wind generates eastward-flowing surface currents with velocities of order 10 cm s\(^{-1}\) outside the inlet (Fig. 10(a)). The orientation of Beaufort Inlet is such that part of this eastward flow is directed into the inlet and is then driven eastward behind Shackleford Banks, through Core Sound and ultimately into Pamlico Sound. Because the eastward wind is roughly parallel to the shoreline in this area, there are few locations where water may pile up against the shore and generate reversing flows at depth. Therefore, near-bottom currents (Fig. 10(b)) closely resemble surface currents, except that they are diminished in magnitude, because of frictional effects.

The wind blowing toward the north generates weaker surface currents (less than 5 cm s\(^{-1}\)) that are aligned with the wind, so are directed primarily onshore (Fig. 10(c)). Water is driven directly into the inlet and then up estuary through the Morehead, Radio Island and Shackleford Channels. Away from the inlet mouth, water piles up along Shackleford Banks and Bogue Banks, driving an offshore flow at depth. However, near the inlet mouth and inside the inlet, the flow is primarily unidirectional over the depth.

Since the FUNDY5 model is linear, its response to winds of different speed and direction follows directly from these examples. A different strength wind will drive proportionally stronger or weaker currents. (Since the surface stress is quadratic in wind speed, this will occur in proportion to the square of the wind speed.) A wind that blows toward any direction other than north or east will generate a circulation that is the vector sum of that driven by the northward component of the wind and that driven by the eastward component of the wind.

Although the wind-driven residual currents are generally weaker than are the tidal residual currents, the wind-driven currents appear strong enough and appear to have areas of sufficient lateral shear to affect larval transport and dispersion in the vicinity of the inlet, particularly under strong wind conditions.

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Figure 10. Steady state circulation near Beaufort Inlet driven by a wind of 7.5 m s⁻¹; (a) surface currents as a result of a wind blowing towards the east; (b) near-bottom currents (1 m above bottom) as a result of wind blowing towards the east; (c) surface currents as a result of wind blowing towards the north; (d) near-bottom currents (1 m above bottom) as a result of wind blowing towards the north.

SIMULATED LARVAL TRANSPORT

Flow fields from the tidal and wind-driven models were used as input to a particle-tracking model (Baptista et al., 1984; Foreman et al., 1992) to examine potential larval transport pathways in the vicinity of Beaufort Inlet. The particle-tracking model was initiated with a rectangular array of 200 particles in one of three locations (west, centre or east) outside the inlet (Fig. 11). Each particle-tracking model run had a duration of 10 M2 tidal cycles (about 124 h), which is a typical time period over which larvae in this stage might remain in the water column (R. Forward, pers. comm.). As described in detail in the following, a large number of particle-tracking model runs were conducted to determine passive transport under mean tidal conditions, the spring-neap cycle, and wind-plus tidally forced conditions.

Results are presented quantitatively in terms of the number of particles crossing ‘virtual nets’ that were placed across the channels immediately inside Beaufort Inlet at the approximate positions of plankton-sampling stations during the March 1996 field experiment (Fig. 11). Net-crossing statistics of model runs with 2000 particles were found to be nearly identical to those from runs with 200 particles, so the smaller number of particles was used for all runs presented in the following. We note that larval sampling conducted during the field study used nets that were relatively small compared with the channel width, so capturing only a tiny fraction of the total fish that passed any location. In trying to represent this with the model and at the same time keeping the particles to a computationally manageable number, we have used nets spanning the entire channel and allowed the particles
Figure 11. West, centre and east initial positions for the 200 particles used in particle model runs. Heavy lines show the positions of 'virtual nets' used to quantify the number of particles passing through the Morehead, Radio Island and Shackleford Channels.

Figure 12. 10-day trajectories for particles released at slack before ebb and transported by $M_2 + M_4 +$ steady residual tidal currents.

to remain in the water column after crossing a net. However, to eliminate the effect of single particles being trapped near a net and crossing it multiple times, particles were only counted the first time that they crossed each net.

Mean tidal forcing

The principal tidal forcing in this region is the semidiurnal $M_2$ constituent. Therefore, we initially simulated larval transport associated with this constituent plus its principal nonlinear overtide ($M_4$) plus the nonlinearily generated tidal residual circulation. The trajectories (after 10 $M_2$ tidal cycles) of all the particles, assuming a slack before ebb release time, are shown in Fig. 12. Particles released west of the inlet are almost exclusively transported up Morehead Channel into the Newport River Estuary and Bogue Sound. Particles released in the centre of the inlet are transported through Morehead Channel, Radio Island Channel and Shackleford Channel. Particles released east of the inlet are primarily transported into Shackleford Channel, although a significant fraction is transported into Radio Island and Morehead Channels. Also, these trajectory lines show a tendency for particles to become entrained in the residual eddy located outside and west of the inlet (Fig. 6).

Although Fig. 12 shows that the trajectory of a particle is dependent on its release point in space, the trajectory also depends on a particle’s release time.

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during the tidal cycle. Therefore, for each of the three spatial initial conditions, 10 model runs were conducted with starting times that were evenly incremented through the tidal cycle. The average number of net crossings as well as the standard deviation were computed for each ensemble of 10 model runs and are reported in Table 2. The average number of net crossings confirms the patterns depicted in Fig. 12. Over half the particles released west of the inlet progress up Morehead Channel, while only a few pass through Radio Island Channel and virtually none moves through Shackleford Channel. Centre release results in nearly equal numbers of particles passing through Morehead and Shackleford Channels. East release results in three-quarters of the particles passing through Shackleford Channel, and approximately 10% passing through Radio Island and Morehead Channels. These results also show that about half the particles did not cross any nets for west and centre releases, although only about a quarter of the particles did not cross a net for the east release. The standard deviations are typically small compared with the means, indicating that there is some sensitivity of the particle track to starting time during the tidal cycle, but that the overall patterns are relatively independent of starting time.

Animations of particle trajectories (not shown) indicate that the principal pathways that particles follow into the inlet are via the near-shore regions where the tidal residual circulation is strong and directed toward the inlet.

Spring-neap effects
The influence of the spring-neap cycle was considered by repeating the runs described already but including the $S_2$ tidal constituent which, in combination with the $M_2$ constituent, generates the spring-neap cycle. In this case, two sets of runs were conducted: one in which the run period was centred around the neap tide and one in which the run period was centred around the spring tide. Statistics for particles not crossing any nets suggest that transport into the estuary during spring (neap) tide increases (decreases) by about 15–20% compared with the $M_2$ tide alone (Table 2). Net-crossing statistics suggest that most of the increase (decrease) in particles entering the estuary occurs in the channel closest to the release points. For example, for a west release, the number of particles entering Morehead Channel increases (decreases) on the spring (neap) tide. The spring tide also appears to be more effective at transporting particles from west to east across the inlet (such as particles passing the Shackleford net that were initially released west of the inlet) than is either the neap or the mean tide. In fact, during the spring tide, exchange between the Morehead and Shackleford Channels is approximately equal in both the east and west directions, whereas west-to-east exchange is essentially nil for the other conditions. Enhanced transport through Shackleford Channel following a west release appears to occur at the expense of transport through Radio Island Channel, suggesting that, on spring tide, particles may overshoot Radio Island Channel and move through Shackleford Channel instead.

Combined wind and tide effects
Subtidal wind-driven circulation combines with tidal residual currents to alter the characteristics of inlet exchange. To investigate this effect, wind-driven FUNDY5 model results were added to the tidal residuals to generate a combined residual current that was used with the mean tidal currents ($M_2 + M_4$) in an

<table>
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<th>Release location</th>
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<th>Stand. dev. (%)</th>
<th>Radio Island Channel Mean (%)</th>
<th>Stand. dev. (%)</th>
<th>Shackleford Channel Mean (%)</th>
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Stand. dev., standard deviation.
additional series of particle-tracking model runs. To account for vertical structure in the wind-driven currents, separate runs were made for particles released at the surface, mid-depth and near bottom (i.e. at the lowest grid point in the FUNDY5 model). Although larvae may be vertically mixed by turbulence while transiting the inlet, no attempt was made to mix particles vertically during the course of particle tracking. Instead, the surface and bottom runs were considered to represent limiting cases for larvae that are able actively to regulate their position in the water column, while the mid-depth run was assumed to be at least partially representative of larvae mixed through the water column.

To account for the range of possible wind conditions, wind speeds of 5 and 10 m s⁻¹ blowing in eight evenly spaced coordinate directions (i.e. N, NE, E, SE, S, SW, W, NW) were considered. As before, the results represent an ensemble of net crossings from particles released at 10 evenly spaced times during the M₂ tidal cycle that are transported passively for 10 M₂ tidal cycles. We note that wind events are rarely constant for

Figure 13. Net-crossing statistics for 10 M₂ tidal cycle long particle-tracking runs using combined wind-driven and mean tidal currents. Each graph shows the average percentage of particles released west of the inlet, at the specified depth, at 10 evenly spaced phases of the tide that either crossed the listed net or crossed none of the nets. The dotted circle in each graph represents net crossings under the mean tidal conditions with no wind. For Radio Island Channel, full scale is 40% of the initially released particles, while full scale is 100% for the other graphs.
Figure 14. Net-crossing statistics for 10 M₂ tidal cycle long particle-tracking runs using combined wind-driven and mean tidal currents. Each graph shows the average percentage of particles released across the centre of the inlet, at the specified depth, at 10 evenly spaced phases of the tide that either crossed the listed net or crossed none of the nets. The dotted circle in each graph represents net crossings under the mean tidal conditions with no wind. For Radio Island Channel, full scale is 40% of the initially released particles, while full scale is 100% for the other graphs.

Particles released west of the inlet
Winds blowing toward the north or north-east enhance total inlet ingress at the surface (shown by a drop in the number of surface particles that did not enter as compared with the case of tides only; Fig. 13) by propelling particles into a favourable position for transport by the flood tide. Wind-driven currents accomplish this by moving particles broadly closer to the inlet or into the near-shore zone, where the net transport is driven by the strong inlet-directed tidal residual currents found there (Fig. 6). Winds in all

five days in this region, so results from this section are probably overestimates of the true impact of wind on larvae ingress.

Figures 13–15 show results for particles released west, across the centre and east of the inlet respectively. The figures represent the percentages of initial particles that crossed a specified net as a function of the direction toward which the wind was blowing. A dotted circle is superimposed on each graph to represent the percentage net crossings in the absence of wind (i.e. as specified in the upper section of Table 2).
Figure 15. Net-crossing statistics for 10 M2 tidal cycle long particle-tracking runs using combined wind-driven and mean tidal currents. Each graph shows the average percentage of particles released east of the inlet, at the specified depth, at 10 evenly spaced phases of the tide that either crossed the listed net or crossed none of the nets. The dotted circle in each graph represents net crossings under the mean tidal conditions with no wind. For Radio Island Channel, full scale is 40% of the initially released particles, while full scale is 100% for the other graphs.

Other directions either blow surface particles away from the inlet or offshore, so decrease ingress. Bottom ingress is enhanced by winds with eastward and southward components. Southward winds transport bottom particles in the onshore direction, while eastward winds transport bottom particles toward the inlet. At mid-depth, enhanced ingress also occurs primarily as a result of southward winds although this enhancement was significant only for the 10 m s⁻¹ wind speed.

Winds generally reduce particle ingress through Morehead Channel. Southward winds are the exception, because the slight increase in total ingress at the bottom appears to be transported up estuary through Morehead Channel. For northward winds, increased total ingress through the inlet plus some of the purely tidal ingress in Morehead Channel are diverted to Radio Island Channel. This appears to be as a result of the up-estuary residual circulation close to the mouth and along the length of Radio Island Channel generated by a north wind (Figs 10(c) and 10(d)). This effect is strong enough that, for the wind speed of 10 m s⁻¹, the number of particles moving through Radio Island
Channel is comparable with that through the much larger Morehead Channel. Eastward winds generate up-estuary currents in Radio Island Channel and Shackleford Channels, so enhancing particle transport through both. Enhanced surface transport into the inlet that occurs with north-eastward wind moves primarily up Shackleford Channel. Eastward and south-eastward winds decrease total ingress at the surface, by blowing particles out of the inlet. However, these wind-driven currents are weaker at the bottom, so are less able to move particles away from the inlet. Instead, it seems that particles are pushed to the east side of the inlet, where they progress up Shackleford Channel. Ingress through Shackleford Channel increases from essentially nil for purely tidal currents to slightly larger than that through Morehead Channel for a wind speed of 10 m s⁻¹.

Partially released across the centre of the inlet

Wind blowing towards the north is particularly effective at enhancing ingress of surface particles as 88% and 94% crossed at least one of the nets for the wind speeds of 5 and 10 m s⁻¹ respectively (compared with only 56% for the purely tidal circulation). The northward wind keeps particles close enough to the inlet such that they can be advected in on a flood tide. Winds blowing in other directions decreased ingress, by creating a surface residual current either offshore or away from the inlet on the downwind side. At the bottom, northward winds also tend to enhance ingress, by driving a current toward the inlet in the inlet centre (Fig. 10(d)). However, a southward wind also enhances ingress, by driving an onshore bottom current both on the eastern and the western sides of the inlet, which brings particles into the near-shore zone where they can then be transported into the inlet by the tidal residual circulation. In addition, an eastward wind enhances bottom ingress, since it generates currents into the inlet that are not strong enough to push particles past the inlet to the east as occurs at the surface. Particles at mid-depth reflect both the surface and bottom behaviours to varying degrees, as ingress is enhanced by both northward and southward winds.

Particles entering at the surface with a northward wind move up estuary through all three inner channels. As before, the largest proportional increase compared with the purely tidal case occurs in Radio Island Channel. Particles entering at the bottom with a southward wind primarly move up Morehead Channel, since this wind generates down-estuary currents in both Radio Island and Shackleford Channels. Particles entering at the bottom with an eastward wind favour up-estuary transport mainly via Shackleford Channel. With this wind, particle numbers are several times greater in Shackleford Channel than is the case in Morehead Channel, as compared with being nearly equally distributed between these channels with purely tidal circulation.

Particles released east of the inlet

Consistent with the other release locations, winds blowing northward are extremely efficient at enhancing surface drifter ingress, as 95% and 97% of the particles crossed at least one of the nets for the wind speeds of 5 and 10 m s⁻¹ respectively. Interestingly, no other wind direction enhances larval ingress at any level in the water column. This is partly because of a relatively high percentage of particles entering the inlet under purely tidal conditions (Table 2). Particle loss from the inlet system was much greater at the surface than it was at depth, presumably because of correspondingly higher wind-driven current speeds. Surface ingress into all three channels was enhanced by the northward wind, while north-westward and southward winds redirect particles from Shackleford Channel into Morehead Channel.

DISCUSSION AND CONCLUSIONS

This study considers barotropic circulation and transport in the vicinity of Beaufort Inlet, NC. Observations indicate that this inlet is weakly stratified at most times (see, for example, Blanton et al., 1999; Churchill et al., 1999), making the barotropic case an important part of a complete understanding of the circulation at this and similar barrier island inlets. We have developed a highly accurate model representation of the inlet geometry and tidal circulation, but simplified the problem by considering steady-state wind responses and restricting particles to remaining passively at the top, middle or bottom of the water column throughout their transit.

Given the high number of scenarios investigated, the assumption of a steady-state wind response was a practical necessity and should not be too unreasonable, owing to the shallow water depths and, hence, rapid spin-up time of the system. However, since weather systems are not normally stationary in this area for as long as 10 M₇ tidal cycles (the duration of the particle-tracking model runs), the computed wind effects on larval transport are probably overestimates of what might actually be expected. Nevertheless, the results provide useful insight into characteristic behaviours of the system for various wind conditions.

An earlier study (Luettich et al., 1998) has shown that simple day-night vertical migratory behaviour causes minimal departure from passive particle transport patterns during March (the primary migratory period in
the inlet part of the SABRE programme), because the day–night cycle and the tidal cycle have nearly the same duration at this time of year. Larvae that move to the bottom at daybreak to avoid visual predation, such that they cease to be transported horizontally, move back up into the water column the following nightfall at approximately the same phase of the tide during which they left. As a result, they follow horizontal paths that are nearly equivalent to those of purely passive particles. Larval catches during the March 1996 field experiments at Beaufort Inlet showed that, in the channels inside the inlet, larvae appeared to be undergoing selective tidal stream transport in which they were mostly present in the water column during nighttime flood tides (Churchill et al., 1999; Forward et al., 1999). However, larval catches in the inlet mouth are not as conclusive and, in general, it is not known where in the system larvae begin this behaviour. By considering passive particle transport, we are only assessing larval transport potential. However, this is a necessary and important step toward understanding more complex larval transport, particularly given the unknowns in larval behaviour outside the inlet.

The vertically integrated ADCIRC tidal model, yielded good agreement with National Ocean Survey tidal data and observations collected during the March 1996 field study. The nonlinearly generated residual current patterns are consistent with available field data and idealized barotropic inlet studies that show offshore-directed flow in the middle of an inlet and flow toward the inlet in the near-shore zones on either side. However, in contrast to an idealized barotropic tidal inlet, the model indicates that a significant residual eddy exists only west of the offshore flow. The structure of the current field suggests that water exiting the inlet would preferentially move offshore, to the west and be re-entrained along the western side of the inlet, while offshore ocean water would be most readily entrained along the eastern side of the inlet. Wind-driven currents generated with the three-dimensional FUN-DY5 model are generally weaker than are the tidal residual currents, although they appear significant enough to affect larval transport in the vicinity of the inlet, particularly during strong wind conditions.

Particle-tracking model runs indicate that, under mean tidal forcing, particles initially released west of the inlet enter the inlet and are transported up-estuary via Morehead Channel. Virtually none crosses the inlet from west to east to move through Shackleford Channel. Particles released across the inlet mouth enter the inlet and are transported up estuary in roughly equal proportions through Morehead and Shackleford Channels. A significantly higher percentage of particles released east of the inlet actually ingress into the system when compared with those released west of and across the inlet mouth. The up-estuary transport of these particles is primarily through Shackleford Channel, although 10–20% of the particles are able to cross the inlet from east to west, so enter through Morehead Channel. Ingress occurs primarily in the near-shore zone that is characterized by tidal residual currents directed toward the inlet. Spring (neap) tide increases (decreases) net particle ingress by 15–20% over the mean tide conditions. During spring tide, exchange between Morehead and Shackleford Channels is approximately equal in both east and west directions.

Particle-tracking model runs that include both wind-driven and the mean tidal circulation indicate that only a narrow range of wind directions significantly enhances particle ingress (compared with ingress as a result of the mean tide alone), while a wide range of wind conditions significantly decrease particle ingress. Northward wind most effectively increases ingress of surface particles from all areas outside the inlet, by pushing these particles either directly into the inlet or into the near-shore zone where tidal currents are favourable for ingress. Inside the inlet, the particle transport path that experiences the greatest increase on a northward wind is Radio Island Channel. This occurs because a northward wind generates an up-estuary circulation near the mouth of Radio Island Channel and along its length. No other wind enhances particle ingress at the surface, mid-depth or bottom from the east side of the inlet. At the bottom of the water column, ingress is enhanced from the centre and the west side of the inlet by southward and eastward winds. Southward wind drives an onshore flow at depth on either side of the inlet, which transports particles onshore and, hence, into the ingress-favourable nearshore zone. An eastward wind moves particles at depth toward the eastern side of the inlet, where they are efficiently transported into the inlet by the tidal circulation. Inside the inlet, particle transport is enhanced in Morehead Channel by southward wind and in Shackleford Channel by eastward wind. Particles at mid-depth reflect both surface and bottom behaviours to varying degrees. In general, their movement is closer to bottom particles for a wind speed of 5 m s⁻¹ and closer to surface particles for a wind speed of 10 m s⁻¹. Stronger wind deepens the surface layer and, therefore, directly influences more of the water column than does lighter wind.

Returning to the observation that nearly an order of magnitude more larvae were collected in Shackleford Channel than in Morehead Channel during the larval transport field experiments in March 1996, our modelling
indicates that, in the absence of wind effects, these larvae must have approached Beaufort Inlet primarily from the east to have attained such a distribution inside the inlet. If the larvae approached Beaufort Inlet in a relatively uniform spatial distribution that spanned both sides of the opening, then the only mechanism for concentrating their numbers in Shackleford Channel vs. Morehead Channel would be an eastward wind. While the climatological mean wind direction in this region is eastward during March, there were no significant eastward winds observed during the March 1996 experiments.

REFERENCES


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